Under the Paperwork Reduction Act of 1995, no persons are required to resp

INFORMATION DISCLOSURE
STATEMENT BY APPLICANT
(Not for submission under 37 CFR 1.99)

Application Number		10692824		
Filing Date		2003-10-23		
First Named Inventor John		Langenfeld		
Art Unit		1643		
Examiner Name Step		nen L. Rawling	_	
Attorney Docket Number		54704.8036.US03(UMD-0072)	_	

					U.S.I	PATENTS			Remove	
Examiner Initial*	Cite No	Patent Number	Kind Code ¹ Issue Date		Date	Name of Patentee or Applicant of cited Document		Pages,Columns,Lines where Relevant Passages or Relevant Figures Appear		
	1									
If you wis	h to a	l dd additional U.S. Pate	nt citatio	n inform	ation pl	ease click the	Add button.		Add	
			U.S.P	ATENT	APPLIC	CATION PUBL	LICATIONS		Remove	
Examiner Initial*	Cite No	Publication Number	Kind Code ¹	Publica Date	tion	Name of Pate of cited Docu	entee or Applicant ment	Releva	Columns,Lines whe int Passages or Rele s Appear	
	1									
If you wisl	h to a	dd additional U.S. Publ	shed Ap	plication	citation	n information p	lease click the Ad	d button	Add	
				FOREIG	SN PAT	ENT DOCUM	ENTS		Remove	
Examiner Initial*	Cite No	Foreign Document Number ³	Country Code ²		Kind Code ⁴	Publication Date	Name of Patente Applicant of cited Document	or v	Pages,Columns,Line where Relevant Passages or Releval Figures Appear	71
	1									
If you wisl	h to a	dd additional Foreign P	atent Do	cument	citation	information pl	ease click the Add	button	Add	
			NON	I-PATE	NT LITE	RATURE DO	CUMENTS		Remove	
Examiner linitials* linclude name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc), date, pages(s), volume-issue number(s), publisher, city and/or country where published.					T					

INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)

Application Number		10692824		
Filing Date		2003-10-23		
First Named Inventor John		Langenfeld		
Art Unit		1643		
Examiner Name	Stept	nen L. Rawling		
Attorney Docket Number		54704.8036.US03(UMD-0072)		

1	Meric et al., "2d 1839 "Iressa", Bull Canoer 2000 87(12):873-876	V
2	Strotnak et al., "Co-administration of probenecid, an inhibitor of a cMOATM/RP-like plasma membrane ATPase, greatly enhanced the efficacy of a new 10-deazasminopterin against human solid furnors in vivo", Clin Cancer Res. 2000 (6)9:3705-3712	Z

If you wish to add additional non-patent literature document citation information please click the Add button Add

EXAMINER SIGNATURE

Examiner Signature Date Considered

*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 809. Draw line through a citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

1 See Kind Codes of USPTIO Patient Documents at sweet USPTIO GOU or MEPE 901.4. ² Eiter office that issued the document, by the Novelets code (WIPO Standard ST3.) ² or Justianese plants columents, be written along the year of the Emperor must preced the series all under of the plant of counters, but will not colument to the control of the propriet and the propriet and the propriet and the propriet and the colument of the propriet and the propriet and the colument of the propriet and the pro

INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)

Application Number		10692824	
Filing Date		2003-10-23	
First Named Inventor	John	Langenfeld	
Art Unit		1643	
Examiner Name	Stept	hen L. Rawling	
Attorney Docket Number		54704.8036.US03(UMD-0072)	

CERTIFICATION STATEMENT

Please see 37 CFR 1.97 and 1.98 to make the appropriate selection(s);

That each item of information contained in the information disclosure statement was first cited in any communication from a foreign patent office in a counterpart foreign application not more than three months prior to the filing of the information disclosure statement. See 37 CFR 197(a)(1).

OR

That no item of information contained in the information disclosure statement was cited in a communication from a foreign patent office in a counterpart foreign application, and, to the knowledge of the person signing the certification after making reasonable inquity, no item of information contained in the information disclosure statement was known to any individual designated in 37 CFR 156(c) more than three months prior to the filing of the information disclosure statement. See 37 CFR 197(s)(c)

- See attached certification statement.
- Fee set forth in 37 CFR 1.17 (p) has been submitted herewith.
- ☐ None

SIGNATURE

A signature of the applicant or representative is required in accordance with CFR 1.33, 10.18. Please see CFR 1.4(d) for the form of the signature.

Signature	/Jane Massey Licata/	Date (YYYY-MM-DD)	2006-07-12

This collection of information is required by 3 T CPR 1.97 and 1.98. The information is required to obtain or retain a benefit by the public which is for life and by the USPTO to process) an application. Confidentially is governed by \$5 U.S. C. 12 and 3 T CPR 1.14. This collection is estimated to take 1 hour to complete, including gathering, preparing and submitting the completed application from the USPTO. The will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, u.S. Operationed of Commence, P. 0. Bot 1450, Alexandria, V.3.251.1450, DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P. 0. Box 1450, Alexandria, V.4.2231.1450.

Privacy Act Statement

The Privacy Act of 1974 (P.L. 93-579) requires that you be given certain information in connection with your submission of the attached form related to a patient application or patient. Accordingly, pursuant to the requirements of the Act, please be advised that. (1) the general authority for the collection of this information is 35 U.S.C. 2(b)(2); (2) famishing of the information solicided is voluntary, and (3) the principal purpose for which the information is used by the U.S. Patient and Trademan Kolfice is to process another examine your submission relation to a patient application or patient. If you do not furnish the requested process another examine your submission relation to the patient application or patient. If you do not furnish the requested the process another examines your submission, which may visually intermediate or for extension or about those when the basic high process another examines your submission, which may visually intermediate or for extension or a submission of the basic high process another examines your submission, which may

The information provided by you in this form will be subject to the following routine uses:

- The information on this form will be treated confidentially to the extent allowed under the Freedom of Information Act (5 U.S.C. 552) and the Privacy Act (5 U.S.C. 552a). Records from this system of records may be disclosed to the Department of Justice to determine whether the Freedom of Information Act requires disclosure of these record s.
 - A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement neodiations.
 - A record in this system of records may be disclosed, as a routine use, to a Member of Congress submitting a request involving an individual, to whom the record partains, when the individual has requested assistance from the Member with respect to the subject matter of the record.
 - A record in this system of records may be disclosed, as a routine use, to a contractor of the Agency having need for the information in order to perform a contract. Recipients of information shall be required to comply with the requirements of the Privacy Act of 1974. as amended, pusuant to 5 U.S.C. 552a(m).
 - A record related to an International Application filed under the Patent Cooperation Treaty in this system of records
 may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant
 to the Patent Cooperation Treaty.
 - A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
- 7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or hisher designe, cuting an inspection of records concluded by GSAs a part of that apency's responsibility to recommend improvements in records management practices and programs, under suthority of 4d U.S.C. 2004 and 2006. Such disclosure shall be made in accordance with the GSA regulations governing inspection of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make determinations about individuals.
- A record from this system of records may be disclosed, as a routine use, to the public after either publication of the application pursuant to 35 U.S. C. 122(b) or issuance of a patent pursuant to 35 U.S. C. 151. Further, a record may be disclosed, subject to the limitations of 37 CFR 1.14, as a routine use, to the public if the record via set float in an application which became abandomed or in which the proceedings were terminated and which application is referenced by either a published application, an application open to public inspections or an issuand patent.
 - A record from this system of records may be disclosed, as a routine use, to a Federal, State, or local law enforcement agency, if the USPTO becomes aware of a violation or potential violation of law or regulation.